

10/584946

Document made available under the Patent Cooperation Treaty (PCT)

International application number: PCT/JP05/000319

International filing date: 06 January 2005 (06.01.2005)

Document type: Certified copy of priority document

Document details: Country/Office: JP
Number: 2004-001311
Filing date: 06 January 2004 (06.01.2004)

Date of receipt at the International Bureau: 03 March 2005 (03.03.2005)

Remark: Priority document submitted or transmitted to the International Bureau in compliance with Rule 17.1(a) or (b)



World Intellectual Property Organization (WIPO) - Geneva, Switzerland
Organisation Mondiale de la Propriété Intellectuelle (OMPI) - Genève, Suisse

04.2.2005

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This is to certify that the annexed is a true copy of the following application as filed with this Office.

出願年月日 2004年 1月 6日
Date of Application:

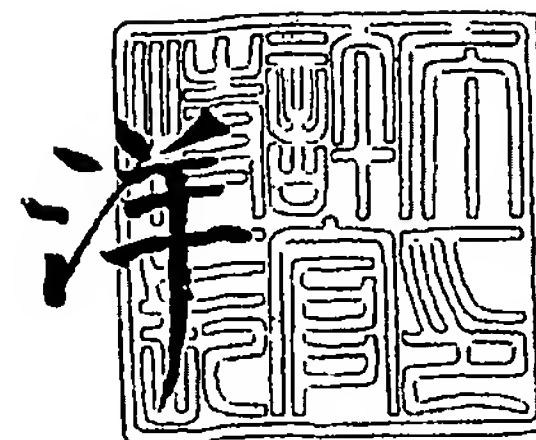
出願番号 特願2004-001311
Application Number:
[ST. 10/C]: [JP2004-001311]

出願人 大正製薬株式会社
Applicant(s):

2005年 1月 5日

特許庁長官
Commissioner,
Japan Patent Office

小川



出証番号 出証特2004-3119372

【書類名】 特許願
【整理番号】 DA-03602
【特記事項】 特許法第36条の2第1項の規定による特許出願
【提出日】 平成16年 1月 6日
【あて先】 特許庁長官殿
【国際特許分類】 C07D419/00
【発明者】
 【住所又は居所】 東京都豊島区高田3丁目24番1号 大正製薬株式会社内
 【氏名】 中里 篤郎
【発明者】
 【住所又は居所】 東京都豊島区高田3丁目24番1号 大正製薬株式会社内
 【氏名】 大久保 武利
【発明者】
 【住所又は居所】 東京都豊島区高田3丁目24番1号 大正製薬株式会社内
 【氏名】 野沢 大
【発明者】
 【住所又は居所】 東京都豊島区高田3丁目24番1号 大正製薬株式会社内
 【氏名】 民田 智子
【発明者】
 【住所又は居所】 ベルギー国、ビールセ、トゥルンハウトセヴェク 30、ヤンセ
 ン ファル マソーティカ エヌ、ヴィー、内
 【氏名】 ケニス、リュド、イー、ジェイ
【特許出願人】
 【識別番号】 000002819
 【氏名又は名称】 大正製薬株式会社
【代理人】
 【識別番号】 100115406
 【弁理士】
 【氏名又は名称】 佐鳥 宗一
【復代理人】
 【識別番号】 100066692
 【弁理士】
 【氏名又は名称】 浅村 皓
【選任した復代理人】
 【識別番号】 100072040
 【弁理士】
 【氏名又は名称】 浅村 肇
【選任した復代理人】
 【識別番号】 100107504
 【弁理士】
 【氏名又は名称】 安藤 克則
【選任した復代理人】
 【識別番号】 100102897
 【弁理士】
 【氏名又は名称】 池田 幸弘
【手数料の表示】
 【予納台帳番号】 002901
 【納付金額】 35,000円
【提出物件の目録】
 【物件名】 外国語特許請求の範囲 1



【物件名】

外国語明細書 1

【物件名】

外国語要約書 1

認定・付加情報

特許出願の番号	特願 2004-001311
受付番号	50400012043
書類名	特許願
担当官	笹川 友子 9482
作成日	平成16年 3月31日

<認定情報・付加情報>

【特許出願人】

【識別番号】	000002819
【住所又は居所】	東京都豊島区高田3丁目24番1号
【氏名又は名称】	大正製薬株式会社

【代理人】

【識別番号】	100115406
【住所又は居所】	東京都豊島区高田3丁目24番1号 大正製薬株式会社 知的財産部

【氏名又は名称】	佐鳥 宗一
----------	-------

【復代理人】

【識別番号】	100066692
【住所又は居所】	東京都千代田区大手町2丁目2番1号 新大手町ビルディング331

【氏名又は名称】	浅村 皓
----------	------

【選任した復代理人】

【識別番号】	100072040
【住所又は居所】	東京都千代田区大手町2丁目2番1号 新大手町ビルディング331

【氏名又は名称】	浅村 肇
----------	------

【選任した復代理人】

【識別番号】	100107504
【住所又は居所】	東京都千代田区大手町2丁目2番1号 新大手町ビルディング331-340

【氏名又は名称】	安藤 克則
----------	-------

【選任した復代理人】

【識別番号】	100102897
【住所又は居所】	東京都千代田区大手町二丁目2番1号 新大手町ビルディング331

【氏名又は名称】	池田 幸弘
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[TITLE OF INVENTION]

PYRROLOPYRIMIDINE AND PYRROLOTRIAZINE DERIVATIVES

[DETAILED DESCRIPTION OF THE INVENTION]

5

[TECHNICAL FIELD]

The present invention relates to a therapeutic agent for diseases in which corticotropin releasing factor (CRF) is considered to be involved, such as depression, anxiety, Alzheimer's disease, Parkinson's disease, Huntington's chorea, eating disorder, hypertension, gastral diseases, drug dependence, cerebral infarction, cerebral ischemia, cerebral edema, cephalic external wound, inflammation, immunity-related diseases, alpecia, irritable bowel syndrome, sleep disorders, epilepsy, dermatitides, schizophrenia, etc.

[DESCRIPTION OF THE PRIOR ART]

15

CRF is a hormone comprising 41 amino acids (Science, 213, 1394-1397, 1981; and J. Neurosci., 7, 88-100, 1987), and it is suggested that CRF plays a core role in biological reactions against stresses (Cell. Mol. Neurobiol., 14, 579-588, 1994; Endocrinol., 132, 723-728, 1994; and Neuroendocrinol. 61, 445-452, 1995). For CRF, there are the following two paths: a path by which CRF acts on peripheral immune system or sympathetic nervous system through hypothalamus-pituitary-adrenal system, and a path by which CRF functions as a neurotransmitter in central nervous system (in Corticotropin Releasing Factor: Basic and Clinical Studies of a Neuropeptide, pp. 29-52, 1990). Intraventricular administration of CRF to hypophysectomized rats and normal rats causes an anxiety-like symptom in both types of rats (Pharmacol. Rev., 43, 425-473, 1991; and Brain Res. Rev., 15, 71-100, 1990). That is, there are suggested the participation of CRF in hypothalamus-pituitary-adrenal system and the pathway by which CRF functions as a neurotransmitter in central nervous system.

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25

The review by Owens and Nemeroff in 1991 summarizes diseases in which CRF is involved (Pharmacol. Rev., 43, 425-474, 1991). That is, CRF is involved in depression, anxiety,

Alzheimer's disease, Parkinson's disease, Huntington's chorea, eating disorder, hypertension, gastrointestinal diseases, drug dependence, inflammation, immunity-related diseases, etc. It has recently been reported that CRF is involved also in epilepsy, cerebral infarction, cerebral ischemia, cerebral edema, and cephalic external wound (Brain Res. 545, 339-342, 1991; Ann. Neurol. 31, 48-498, 1992; Dev. Brain Res. 91, 245-251, 1996; and Brain Res. 744, 166-170, 1997). Accordingly, antagonists against CRF receptors are useful as therapeutic agents for the diseases described above.

[PROBLEM(S) TO BE SOLVED BY INVENTION]

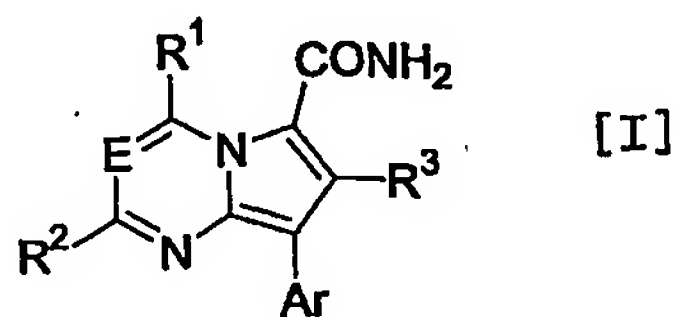
10 An object of the present invention is to provide an antagonist against CRF receptors which is effective as a therapeutic or prophylactic agent for diseases in which CRF is considered to be involved, such as depression, anxiety, Alzheimer's disease, Parkinson's disease, Huntington's chorea, eating disorder, hypertension, gastral diseases, drug dependence, epilepsy, cerebral infarction, cerebral ischemia, cerebral edema, cephalic external wound, inflammation, 15 immunity-related diseases, alpecia, irritable bowel syndrome, sleep disorders, epilepsy, dermatitides, schizophrenia, etc.

[MEANS FOR SOLVING PROBLEM]

The present inventors earnestly investigated pyrrolopyrimidines or pyrrolotriazines 20 substituted with a carbamoyl group that have a high affinity for CRF receptors, whereby the present invention has been accomplished.

The present invention is pyrrolopyrimidine or pyrrolotriazine derivatives substituted with a carbamoyl group explained below.

A pyrrolopyrimidine or pyrrolotriazine derivative substituted with a carbamoyl group 25 represented by the following formula [I]:



(wherein E is N or CH;

R^1 is $-OR^4$, $-S(O)_lR^4$ or $-NR^4R^5$;

R^4 and R^5 are the same or different, and independently hydrogen, C_{1-9} alkyl, C_{3-7} cycloalkyl, C_{3-7} cycloalkyl- C_{1-6} alkyl, di(C_{3-7} cycloalkyl)- C_{1-6} alkyl, C_{1-6} alkoxy- C_{1-6} alkyl, di(C_{1-6} alkoxy)- C_{1-6} alkyl, hydroxy- C_{1-6} alkyl, cyano- C_{1-6} alkyl, carbamoyl- C_{1-6} alkyl or di(C_{1-6} alkyl)amino- C_{2-6} alkyl; or R^4 and R^5 are taken together to form $-(CH_2)_m-A-(CH_2)_n-$ wherein A is methylene, oxygen, sulfur, NR^6 or CHR^7 , m is an integer selected from 1, 2, 3 and 4, n is an integer selected from 0, 1 and 2, wherein R^6 is hydrogen, C_{1-6} alkyl or C_{3-7} cycloalkyl, R^7 is hydroxy, hydroxy- C_{1-6} alkyl, cyano or cyano- C_{1-6} alkyl;

R^2 is hydrogen or C_{1-6} alkyl;

R^3 is hydrogen or C_{1-6} alkyl;

l is an integer selected from 0, 1 and 2;

Ar is aryl or heteroaryl which aryl or heteroaryl is unsubstituted or substituted with 1 or more substituents, which are the same or different, selected from the group consisting of halogen, C_{1-6} alkyl, C_{3-7} cycloalkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{1-6} alkoxy, C_{1-6} alkylthio, cyano, trifluoromethyl, trifluoromethoxy, difluoromethoxy, fluoromethoxy and $-N(R^8)R^9$, wherein R^8 and R^9 are the same or different, and independently are hydrogen or C_{1-6} alkyl), individual isomers thereof or racemic or non-racemic mixtures of isomers thereof, or pharmaceutically acceptable salts and hydrates thereof.

The terms used in the present specification have the following meanings.

The term " C_{1-9} alkyl" means a straight chain or branched chain alkyl group of 1 to 9 carbon atoms, such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, *tert*-butyl, *sec*-butyl, pentyl, isopentyl, 1-methylbutyl, hexyl, isohexyl, 1-ethylpropyl, 1-ethylbutyl, 1,3-dimethylbutyl, 1-propylbutyl, 1-propylpentyl, 1-butylpentyl or the like.

The term " C_{3-7} cycloalkyl" means a cyclic alkyl group of 3 to 7 carbon atoms, such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl or the like.

The term " C_{3-7} cycloalkyl- C_{1-6} alkyl" means a substituted C_{1-6} alkyl group having the

above-mentioned C_{3-7} cycloalkyl as the substituent, such as cyclopropylmethyl, 1-cyclopropylethyl, 1-cyclobutylethyl, 1-cyclopentylethyl, 2-cyclopropylethyl, 2-cyclobutyl-ethyl, 2-cyclopentylethyl, 1-cyclopropyl-propyl, 1-cyclobutyl-propyl, 1-cyclopentyl-propyl, 1-cyclopropylmethyl-propyl, 1-cyclopropylmethyl-butyl or the like.

- 5 The term "di(C_{3-7} cycloalkyl)- C_{1-6} alkyl" means a substituted C_{1-6} alkyl group having two above-mentioned C_{3-7} cycloalkyl groups as the substituents, such as di(cyclopropyl)methyl, di(cyclobutyl)methyl, di(cyclopentyl)methyl or the like.

- 10 The term " C_{1-6} alkoxy" means a straight chain or branched chain alkoxy group of 1 to 6 carbon atoms, such as methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutyloxy, pentyloxy, isopentyloxy or the like.

The term " C_{1-6} alkoxy- C_{1-6} alkyl" means a substituted C_{1-6} alkyl group having the above-mentioned C_{1-6} alkoxy group as the substituent, such as methoxymethyl, 2-methoxyethyl, 2-ethoxyethyl, 1-methoxymethyl-propyl, 1-methoxymethyl-butyl or the like.

- 15 The term "di(C_{1-6} alkoxy)- C_{1-6} alkyl" means a substituted C_{1-6} alkyl group having two above-mentioned C_{1-6} alkoxy groups as the substituents, such as 2,3-di(methoxy)propyl, 2-methoxy-1-methoxymethyl-ethyl, 2,4-(diethoxy)pentyl or the like.

- 20 The term "hydroxy- C_{1-6} alkyl" means a substituted C_{1-6} alkyl group having a hydroxy group, such as hydroxymethyl, 1-hydroxyethyl, 2-hydroxyethyl, 1-hydroxypropyl, 2-hydroxypropyl, 3-hydroxypropyl, 4-hydroxybutyl, 5-hydroxypentyl, 1-hydroxymethyl-propyl, 1-hydroxymethyl-butyl, 1-hydroxymethyl-3-methyl-butyl or the like.

The term "cyano- C_{1-6} alkyl" means a substituted C_{1-6} alkyl group having a cyano group, such as cyanomethyl, 1-cyanoethyl, 2-cyanoethyl, 1-cyanopropyl, 1-cyanobutyl, 5-cyanopentyl, 2-cyano-1-ethyl-ethyl, 1-cyanomethyl-butyl, 1-cyano-3-methyl-butyl, 1-cyanomethyl-3-methyl-butyl or the like.

- 25 The term "carbamoyl- C_{1-6} alkyl" means a substituted C_{1-6} alkyl group having a carbamoyl group, such as carbamoylmethyl, 1-carbamoylethyl, 2-carbamoylethyl, 1-carbamoylpropyl, 1-carbamoylbutyl, 5-carbamoylpentyl, 1-carbamoyl-3-methyl-butyl, 1-carbamoylmethyl-butyl, 1-carbamoylmethyl-propyl, 1-carbamoylmethyl-3-methyl-butyl or the like.

The term "di(C₁₋₆alkyl)amino" means an amino group having two above-mentioned C₁₋₆alkyl groups, such as dimethylamino, diethylamino, dipropylamino or the like.

The term "di(C₁₋₆alkyl)amino-C₂₋₆alkyl" means a substituted C₂₋₆alkyl group having the above-mentioned di(C₁₋₆alkyl)amino group, such as 2-dimethylaminoethyl, 3-dimethylaminopropyl or the like.

The term "aryl" means a monocyclic or bicyclic group of 6 to 12 ring carbon atoms having at least one aromatic ring, such as phenyl, naphthyl, or the like.

The term "heteroaryl" means a monocyclic or bicyclic group of 5 to 12 ring atoms having at least one aromatic ring having in its ring 1 to 4 atoms which may be the same or different and are selected from nitrogen, oxygen and sulfur, such as pyridyl, pyrimidinyl, imidazolyl, quinolyl, indolyl, benzofuranyl, quinoxaliny, benzo[1,2,5]thiadiazolyl, benzo[1,2,5]oxadiazolyl or the like.

The term "halogen" means fluorine, chlorine, bromine or iodine atom.

The term "C₂₋₆alkenyl" means a straight chain or branched chain alkenyl group of 2 to 6 carbon atoms, such as vinyl, isopropenyl, allyl or the like.

The term "C₂₋₆alkynyl" means a straight chain or branched chain alkynyl group of 2 to 6 carbon atoms, such as ethynyl, prop-1-ynyl, prop-2-ynyl or the like.

The term "C₁₋₆alkylthio" means a straight chain or branched chain alkylthio group of 1 to 6 carbon atoms, such as methylthio, ethylthio, propylthio or the like.

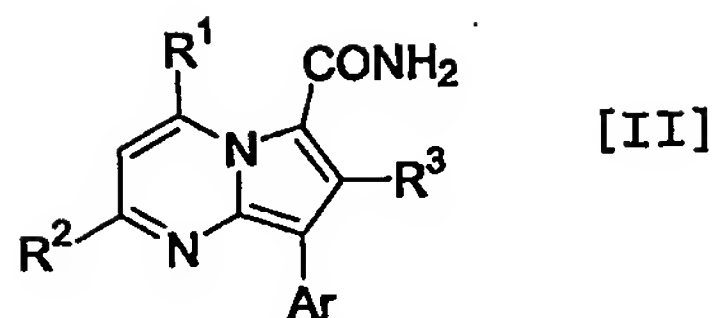
The phrase "aryl or heteroaryl which aryl or heteroaryl is unsubstituted or substituted with 1 or more substituents, which are the same or different, selected from the group consisting of halogen, C₁₋₆alkyl, C₃₋₇cycloalkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₁₋₆alkoxy, C₁₋₆alkylthio, cyano, trifluoromethyl, trifluoromethoxy, difluoromethoxy, fluoromethoxy and -N(R⁷)R⁸, wherein R⁷ and R⁸ are the same or different, and independently are hydrogen or C₁₋₆alkyl" includes, for example, 2,4-dimethylphenyl, 2,6-dimethylphenyl, 2,4-dibromophenyl, 2-bromo-4-isopropylphenyl, 2,4-dichlorophenyl, 2,6-dichlorophenyl, 2-chloro-4-trifluoromethylphenyl, 4-methoxy-2-methylphenyl, 2-chloro-4-trifluoromethoxyphenyl, 4-isopropyl-2-methylthiophenyl, 2,4,6-trimethylphenyl, 4-bromo-2,6-dimethylphenyl, 4-bromo-2,6-diethylphenyl, 4-chloro-2,6-

dimethylphenyl, 2,4,6-tribromophenyl, 2,4,5-tribromophenyl, 2,4,6-trichlorophenyl, 2,4,5-trichlorophenyl, 4-bromo-2,6-dichlorophenyl, 6-chloro-2,4-dibromophenyl, 2,4-dibromo-6-fluorophenyl, 2,4-dibromo-6-methylphenyl, 2,4-dibromo-6-methoxyphenyl, 2,4-dibromo-6-methylthiophenyl, 2,6-dibromo-4-isopropylphenyl, 2,6-dibromo-4-trifluoromethylphenyl, 2-bromo-4-trifluoromethylphenyl, 4-bromo-2-chlorophenyl, 2-bromo-4-chlorophenyl, 4-bromo-2-methylphenyl, 4-chloro-2-methylphenyl, 2,4-dimethoxyphenyl, 2,6-dimethyl-4-methoxyphenyl, 4-chloro-2,6-dibromophenyl, 4-bromo-2,6-difluorophenyl, 2,6-dichloro-4-trifluoromethylphenyl, 2,6-dichloro-4-trifluoromethoxyphenyl, 2,6-dibromo-4-trifluoromethoxyphenyl, 2-chloro-4,6-dimethylphenyl, 2-bromo-4,6-dimethoxyphenyl, 2-bromo-4-isopropyl-6-methoxyphenyl, 2,4-dimethoxy-6-methylphenyl, 6-dimethylamino-4-methylpyridin-3-yl, 2-chloro-6-trifluoromethylpyridin-3-yl, 2-chloro-6-trifluoromethoxypyridin-3-yl, 2-chloro-6-methoxypyridin-3-yl, 6-methoxy-2-trifluoromethylpyridin-3-yl, 2-chloro-6-difluoromethylpyridin-3-yl, 6-methoxy-2-methylpyridin-3-yl, 2,6-dimethoxypyridin-3-yl, 4,6-dimethyl-2-trifluoromethylpyrimidin-5-yl, 2-dimethylamino-6-methylpyridin-3-yl.

15 The "pharmaceutically acceptable salts" in the present invention include, for example, salts with an inorganic acid such as sulfuric acid, hydrochloric acid, hydrobromic acid, phosphoric acid, nitric acid or the like; salts with an organic acid such as acetic acid, oxalic acid, lactic acid, tartaric acid, fumaric acid, maleic acid, citric acid, benzenesulfonic acid, methanesulfonic acid, *p*-toluenesulfonic acid, benzoic acid, camphorsulfonic acid, ethanesulfonic acid, glucoheptonic acid, gluconic acid, glutamic acid, glycolic acid, malic acid, malonic acid, mandelic acid, galactaric acid, naphthalene-2-sulfonic acid or the like; salts with one or more metal ions such as lithium ion, sodium ion, potassium ion, calcium ion, magnesium ion, zinc ion, aluminium ion or the like; salts with an amine such as ammonia, arginine, lysine, piperazine, choline, diethylamine, 4-phenylcyclohexylamine, 2-aminoethanol, benzathine or the like.

25 In a compound of the present invention, isomers such as diastereomers, enantiomers, geometricisomers and tautomeric forms may exist. The compound of the present invention includes the individual isomers and the racemic and non-racemic mixtures of the isomers.

Preferable examples of the compound of the present invention are as follows.

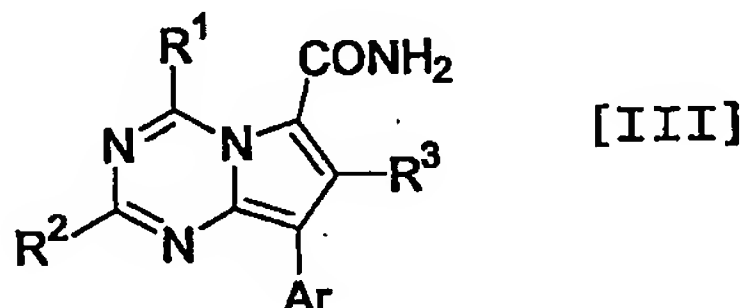


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That is preferable are compounds of the formula [II] in which R^1 , R^2 , R^3 and Ar are as defined in claim 1. More preferable are compounds of the formula [II], wherein R^1 is $-NR^4R^5$; R^4 and R^5 are the same or different, and independently hydrogen, C_{1-9} alkyl, C_{3-7} cycloalkyl, C_{3-7} cycloalkyl- C_{1-6} alkyl, di(C_{3-7} cycloalkyl)- C_{1-6} alkyl, C_{1-6} alkoxy- C_{1-6} alkyl, di(C_{1-6} alkoxy)- C_{1-6} alkyl;

10 R^2 is C_{1-6} alkyl; Ar is phenyl which phenyl is substituted with two or three substituents, which are the same or different, selected from the group consisting of halogen, C_{1-3} alkyl, C_{1-3} alkoxy, C_{1-3} alkylthio, trifluoromethyl, trifluoromethoxy and $-N(R^8)R^9$ (wherein R^8 and R^9 are the same or different, and independently are hydrogen or C_{1-3} alkyl); R^3 is as defined in claim 1.

15



Other preferable are compounds of the formula [III] in which R^1 , R^2 , R^3 and Ar are as defined in claim 1. More preferable are compounds of the formula [III], wherein R^1 is $-NR^4R^5$;

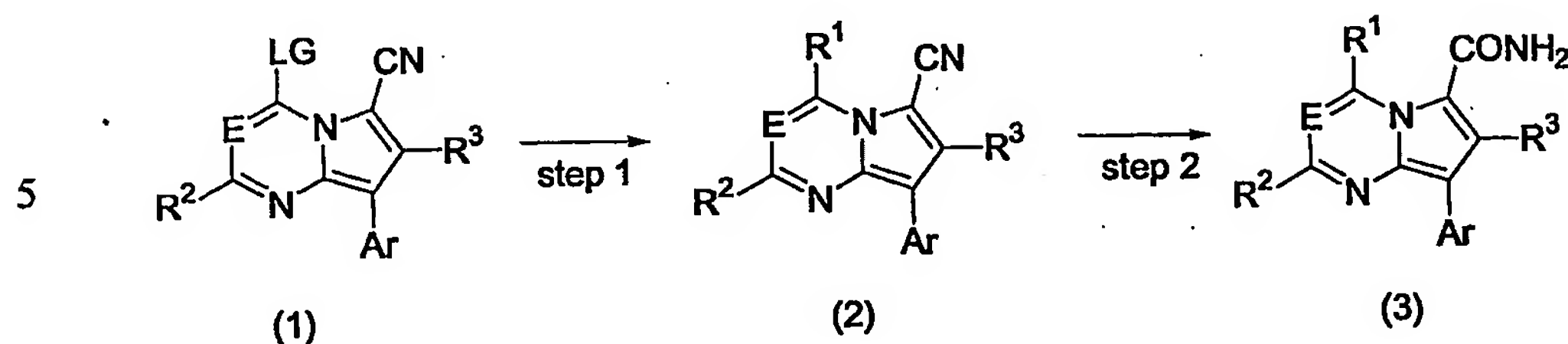
20 R^4 and R^5 are the same or different, and independently hydrogen, C_{1-9} alkyl, C_{3-7} cycloalkyl, C_{3-7} cycloalkyl- C_{1-6} alkyl, di(C_{3-7} cycloalkyl)- C_{1-6} alkyl, C_{1-6} alkoxy- C_{1-6} alkyl, di(C_{1-6} alkoxy)- C_{1-6} alkyl; R^2 is C_{1-6} alkyl; Ar is phenyl which phenyl is substituted with two or three substituents, which are the same or different, selected from the group consisting of halogen, C_{1-3} alkyl, C_{1-3} alkoxy, C_{1-3} alkylthio, trifluoromethyl, trifluoromethoxy and $-N(R^8)R^9$ (wherein R^8 and R^9 are the same or

25 different, and independently are hydrogen or C_{1-3} alkyl); R^3 is as defined in claim 1.

The compound of the formula [I] can be produced, for example, by the process shown in the following reaction scheme 1 (in the following reaction scheme, R^1 , R^2 , R^3 and Ar are as defined above, LG is chloro, bromo, iodo, methanesulfonyloxy, benzenesulfonyloxy,

toluenesulfonyloxy or trifluoromethanesulfonyloxy group, R^a is C_{1-6} alkyl or benzyl, p is 1 or 2).

Reaction Scheme 1



Step 1:

Compound (2), can be obtained by reacting Compound (1) with the corresponding amine
 10 in an inert solvent in the presence or absence of a base. Herein, the base includes, for example, amines such as triethylamine, *N,N*-diisopropylethylamine, pyridine and the like; inorganic bases such as sodium carbonate, potassium carbonate, sodium hydrogencarbonate, potassium hydrogencarbonate, potassium hydroxide, sodium hydroxide, lithium hydroxide, barium hydroxide, sodium hydride and the like; metal alcoholates such as sodium methoxide, sodium
 15 ethoxide, potassium *tert*-butoxide and the like; metal amides such as sodium amide, lithium diisopropylamide and the like; and Grignard reagents such as methylmagnesium bromide and the like. The inert solvent includes, for example, alcohols such as methanol, ethanol, isopropyl alcohol, ethylene glycol and the like; ethers such as diethyl ether, tetrahydrofuran, 1,4-dioxane, 1,2-dimethoxyethane and the like; hydrocarbons such as benzene, toluene, xylene and the like;
 20 amides such as *N,N*-dimethylformamide, *N*-methylpyrrolidone, *N,N*-dimethylacetamide and the like; acetonitrile; dichloromethane; chloroform; dimethyl sulfoxide; pyridine; water; and mixtures of solvents selected from these inert solvents.

Step 2:

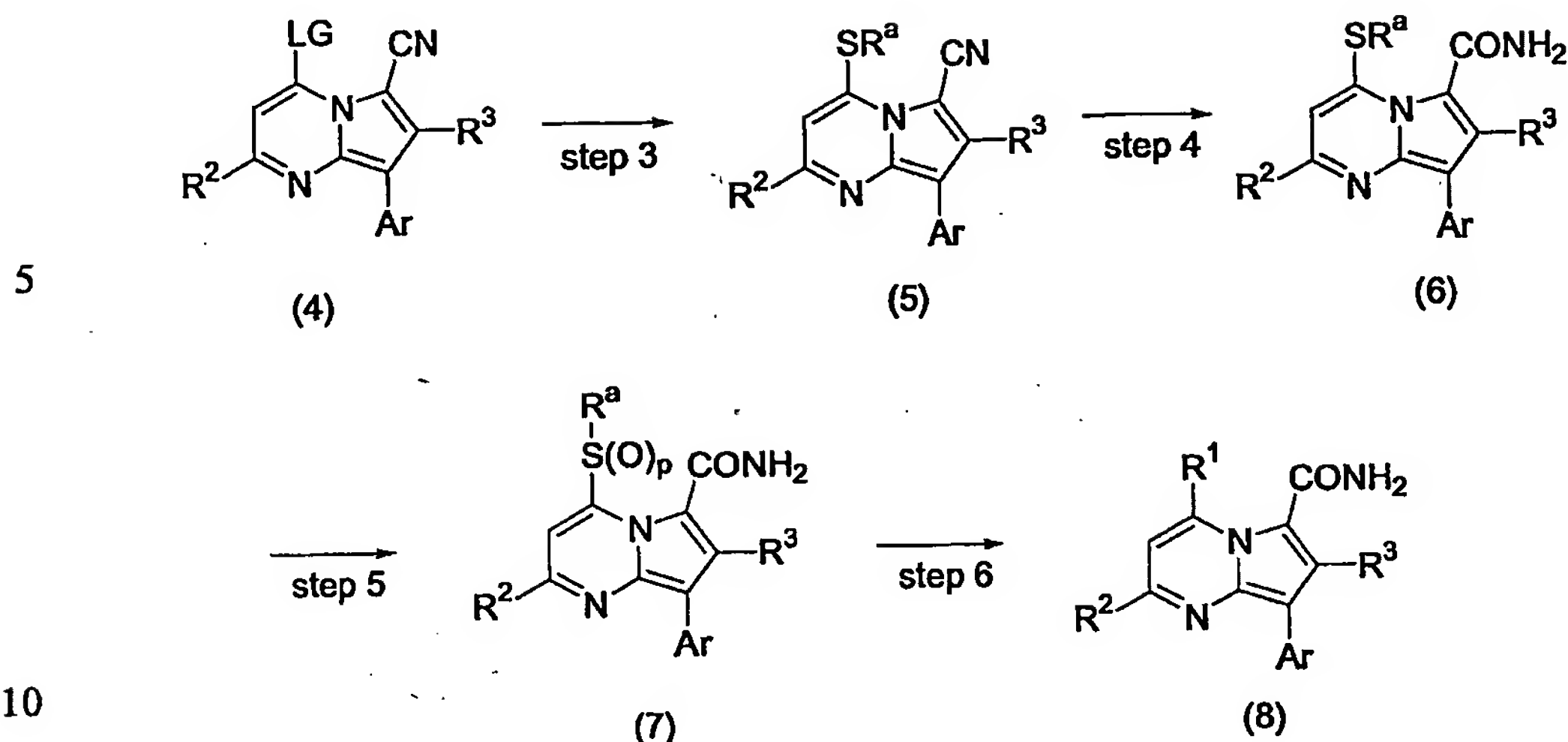
25 Conversion of a cyano group in Compound (2) into a carbamoyl group can be achieved in the presence of an acid or a base in the presence or absence of an inert solvent. When R^1 has a cyano group, the cyano group can be converted into a carbamoyl group at the same time. Herein, the acid includes inorganic acids such as sulfuric acid, hydrochloric acid, hydrobromic acid,

phosphoric acid, polyphosphoric acid nitric acid and the like; organic acids such as benzenesulfonic acid, toluenesulfonic acid and the like. The base includes inorganic bases such as lithium hydroxide, sodium hydroxide, potassium hydroxide, calcium hydroxide, magnesium hydroxide, zinc hydroxide, aluminium hydroxide and the like. The inert solvent includes, for example, alcohols such as methanol, ethanol, isopropyl alcohol, *tert*-butyl alcohol, ethylene glycol and the like; ethers such as diethyl ether, tetrahydrofuran, 1,4-dioxane, 1,2-dimethoxyethane and the like; hydrocarbons such as benzene, toluene and the like; amides such as *N,N*-dimethylformamide, *N*-methylpyrrolidone, *N,N*-dimethylacetamide and the like; acetonitrile; dichloromethane; chloroform; dimethyl sulfoxide; pyridine; water; and mixtures of solvents selected from these inert solvents.

The compound of the present invention can be converted to a salt with an acid in an inert solvent. The acid includes inorganic acids such as sulfuric acid, hydrochloric acid, hydrobromic acid, phosphoric acid, nitric acid and the like; organic acids such as acetic acid, oxalic acid, lactic acid, tartaric acid, fumaric acid, maleic acid, citric acid, benzenesulfonic acid, methanesulfonic acid, *p*-toluenesulfonic acid, benzoic acid, camphorsulfonic acid, ethanesulfonic acid, glucoheptonic acid, gluconic acid, glutamic acid, glycolic acid, malic acid, malonic acid, mandelic acid, galactaric acid, naphthalene-2-sulfonic acid and the like.

The inert solvent includes, for example, alcohols such as methanol, ethanol, isopropyl alcohol, ethylene glycol and the like; ethers such as diethyl ether, tetrahydrofuran, 1,4-dioxane, 1,2-dimethoxyethane and the like; hydrocarbons such as benzene, toluene and the like; amides such as *N,N*-dimethylformamide, *N*-methylpyrrolidone, *N,N*-dimethylacetamide and the like; esters such as ethyl acetate, ethyl formate and the like; ketones such as acetone, methylethylketone and the like; acetonitrile; dichloromethane; chloroform; dimethyl sulfoxide; pyridine; water; and mixtures of solvents selected from these inert solvents.

Reaction Scheme 2



Step 3:

Conversion of Compound (4) into Compound (5) can be carried out by treatment of (4) with thiourea in an inert solvent and followed by reacting with an alkylating reagent in the presence or absence of a base in an inert solvent. The alkylating reagent includes conventional

15 alkylating reagents such as methyl iodide, methyl bromide, dimethyl sulfate, ethyl iodide, ethyl bromide, diethyl sulfate, benzyl chloride, benzyl bromide and the like. The base includes amines such as triethylamine, *N,N*-diisopropylethylamine, pyridine and the like; inorganic bases such as sodium carbonate, potassium carbonate, sodium hydrogencarbonate, potassium

20 hydrogencarbonate, potassium hydroxide, sodium hydroxide, lithium hydroxide, barium hydroxide, sodium hydride and the like; metal alcoholates such as sodium methoxide, sodium ethoxide, potassium *tert*-butoxide and the like; metal amides such as sodium amide, lithium diisopropylamide and the like; and Grignard reagents such as methylmagnesium bromide and the like. The inert solvent includes, for example, alcohols such as methanol, ethanol, isopropyl

25 alcohol, ethylene glycol and the like; ethers such as diethyl ether, tetrahydrofuran, 1,4-dioxane, 1,2-dimethoxyethane and the like; hydrocarbons such as benzene, toluene, xylene and the like; amides such as *N,N*-dimethylformamide, *N*-methylpyrrolidone, *N,N*-dimethylacetamide and the like; acetonitrile; dichloromethane; chloroform; dimethyl sulfoxide; pyridine; water; and mixtures of solvents selected from these inert solvents.

Step 4:

Conversion of Compound (5) into Compound (6) can be achieved in the same manner as step 2.

5 Step 5:

Conversion of Compound (6) into Compound (7) can be carried out by reacting Compound (6) with an oxidizing reagent in an inert solvent. Herein, the oxidizing reagent includes conventional oxidizing reagents to oxidize a sulfide group such as peroxyacetic acid, hydrogen peroxide, 3-chloroperoxybenzoic acid, Oxone, sodium periodate, sodium perborate and
10 the like. The inert solvent includes, for example, alcohols such as methanol, ethanol, isopropyl alcohol, ethylene glycol and the like; ethers such as diethyl ether, tetrahydrofuran, 1,4-dioxane, 1,2-dimethoxyethane and the like; hydrocarbons such as benzene, toluene, xylene and the like; amides such as *N,N*-dimethylformamide, *N*-methylpyrrolidone, *N,N*-dimethylacetamide and the like; acetonitrile; dichloromethane; chloroform; dimethyl sulfoxide; pyridine; water; and mixtures
15 of solvents selected from these inert solvents.

Step 6:

Conversion of Compound (7) into Compound (8) can be carried out in the same manner as step 1.

20

The compound of the present invention is useful as a therapeutic or prophylactic agent for diseases in which CRF is considered to be involved. For this purpose, the compound of the present invention can be formulated into tablets, pills, capsules, granules, powders, solutions, emulsions, suspensions, injections and the like by a conventional preparation technique by adding
25 conventional fillers, binders, disintegrators, pH-adjusting agents, solvents, etc.

The compound of the present invention can be administered to an adult patient in a dose of 0.1 to 500 mg per day in one portion or several portions orally or parenterally. The dose can be properly increased or decreased depending on the kind of a disease and the age, body weight

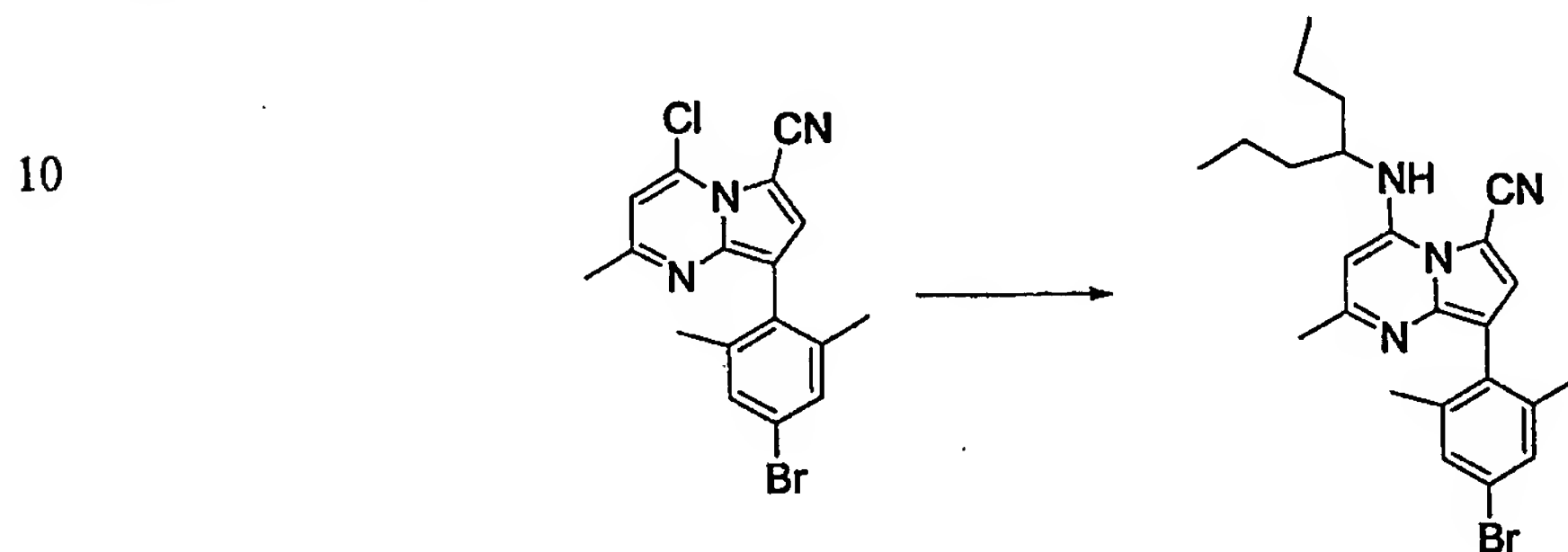
and symptom of a patient.

[EMBODIMENTS OF THE INVENTION]

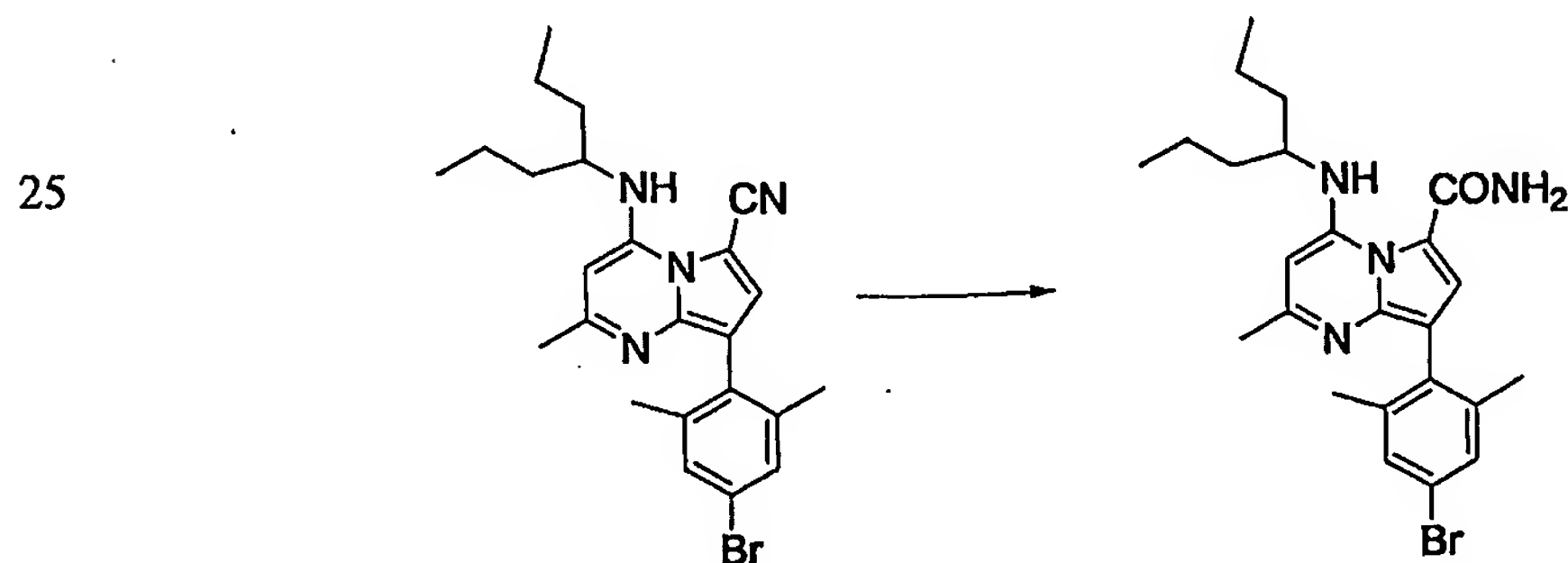
The present invention is concretely explained with reference to the following examples and test example, but is not limited thereto.

Example 1

Synthesis of 8-(4-bromo-2,6-dimethyl-phenyl)-2-methyl-4-(1-propyl-butylamino)-pyrrolo[1,2-*a*]pyrimidine-6-carboxylic acid amide hydrochloride (compound 1-001)



15 (1) A mixture of 8-(4-bromo-2,6-dimethyl-phenyl)-4-chloro-2-methyl-pyrrolo[1,2-*a*]pyrimidine-6-carbonitrile (30.0 g), 1-propyl-butylamine (18.5 g), *N,N*-diisopropylethylamine (15.5 g) in ethanol (90 mL) was heated at reflux for 2 h. The reaction mixture was cooled to room temperature, poured into a saturated aqueous sodium hydrogencarbonate, and then extracted with ethyl acetate. The organic layer was washed with brine, dried over anhydrous sodium sulfate and
20 filtered. The filtrate was concentrated under reduced pressure to give a solid. The solid was washed with diisopropylether to give 8-(4-bromo-2,6-dimethyl-phenyl)-2-methyl-4-(1-propyl-butylamino)-pyrrolo[1,2-*a*]pyrimidine-6-carbonitrile (27.0 g).



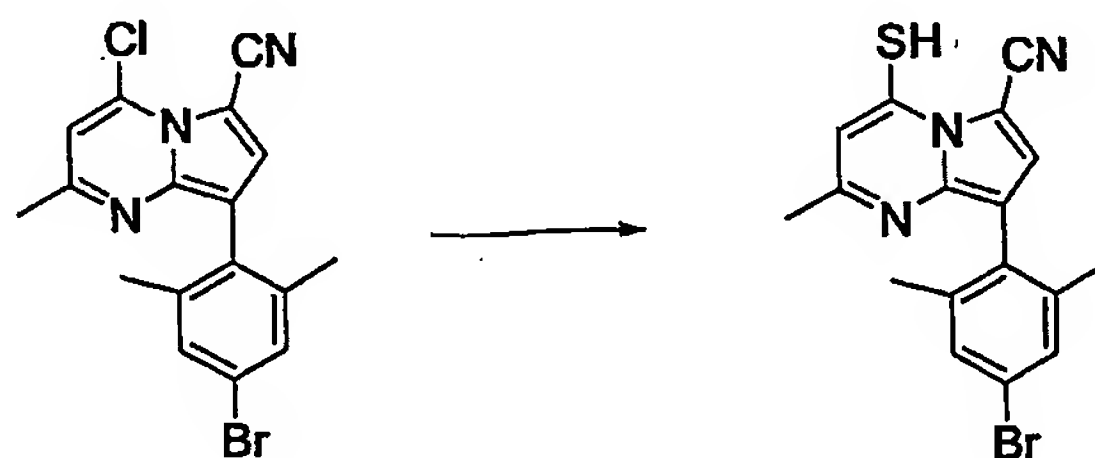
(2) 8-(4-Bromo-2,6-dimethyl-phenyl)-2-methyl-4-(1-propyl-butylamino)-pyrrolo[1,2-*a*]pyrimidine-6-carbonitrile (10.0 g) was added into conc. H₂SO₄ (50 mL) and heated for 55 °C for 5 hours. The reaction mixture was cooled to room temperature, poured into ice-water and then a saturated aqueous sodium hydrogencarbonate was added to make the aqueous mixture alkaline (pH = 8) and extracted with ethyl acetate. The organic layer was washed with brine, dried over anhydrous sodium sulfate and filtered. The filtrate was concentrated under reduced pressure and the residue was purified by a silica gel column chromatography (silica gel: Wako Gel (C200), eluent: hexane / ethyl acetate / chloroform = 10 : 3 : 1) to give a solid. The solid was recrystallized from ethyl acetate to provide 8-(4-bromo-2,6-dimethyl-phenyl)-2-methyl-4-(1-propyl-butylamino)-pyrrolo[1,2-*a*]pyrimidine-6-carboxylic acid amide (5.8 g).

(3) To a suspension of 8-(4-bromo-2,6-dimethyl-phenyl)-2-methyl-4-(1-propyl-butylamino)-pyrrolo[1,2-*a*]pyrimidine-6-carboxylic acid amide (5.8 g) in ethanol (30 mL) was added 4 M HCl / ethyl acetate (3.7 mL) in an ice-cooling bath. The resulting solution was concentrated under reduced pressure. The residue was crystallized from ethyl acetate to give the title compound.

Table 1 and table 2 list the compound obtained in Example 1 and compounds obtained by the similar procedure as in Example 1.

Example 2

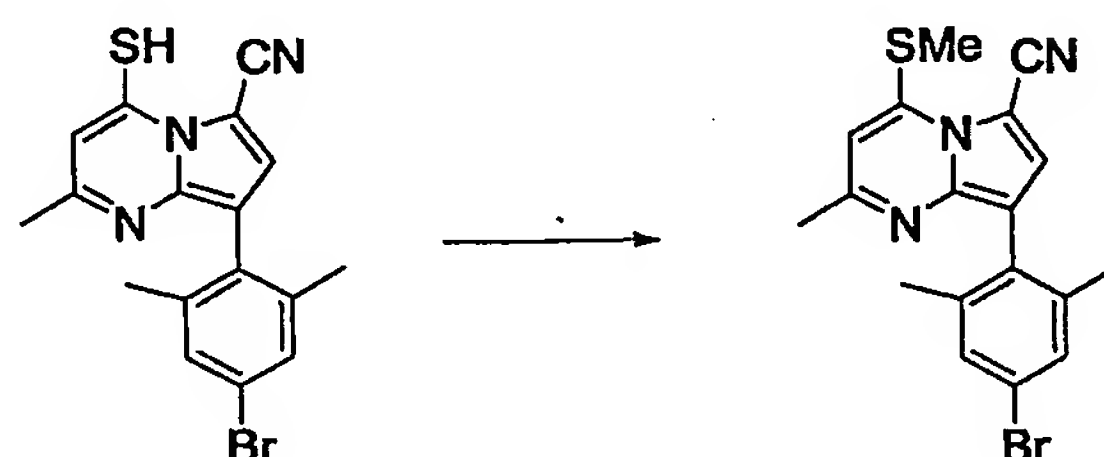
Synthesis of 8-(4-bromo-2,6-dimethyl-phenyl)-2-methyl-4-(*N,N*-dipropylamino)-pyrrolo[1,2-*a*]pyrimidine-6-carboxylic acid amide (compound 1-020)



(1) A mixture of 8-(4-bromo-2,6-dimethyl-phenyl)-4-chloro-2-methyl-pyrrolo[1,2-*a*]pyrimidine-6-carbonitrile (7.50 g), thiourea (7.11 g) in ethanol (50 mL) was heated at reflux for

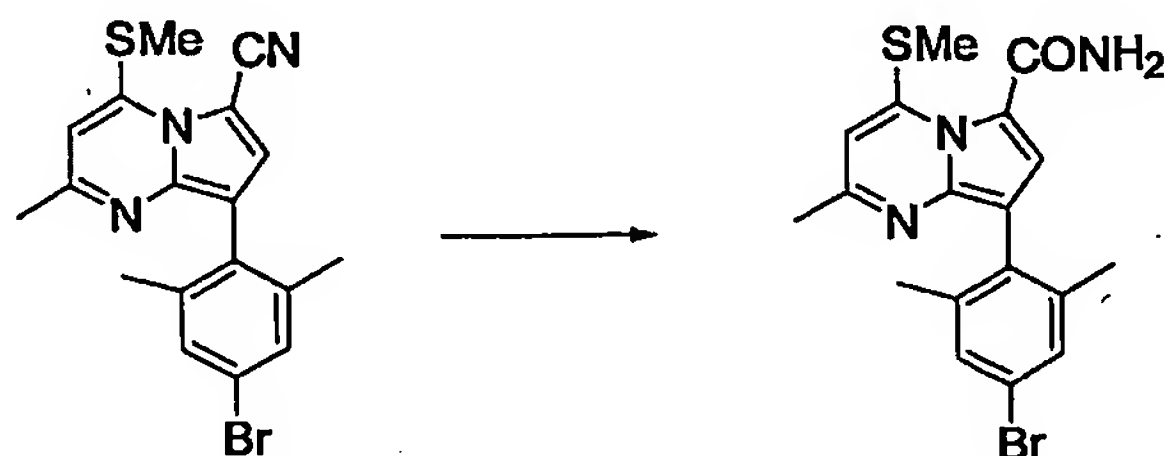
2 h. The reaction mixture was cooled to room temperature, poured into 0.5 M NaOH aqueous solution, stirred for 1 hour and extracted with ethyl acetate. The organic layer was washed with brine, dried over anhydrous sodium sulfate and filtered. The filtrate was concentrated under reduced pressure and the residue was purified by a silica gel column chromatography (silica gel: Wako Gel (C200), eluent: chloroform / methanol = 10 : 1) to give 8-(4-bromo-2,6-dimethyl-phenyl)-4-mercapto-2-methyl-pyrrolo[1,2-*a*]pyrimidine-6-carbonitrile (7.52 g).

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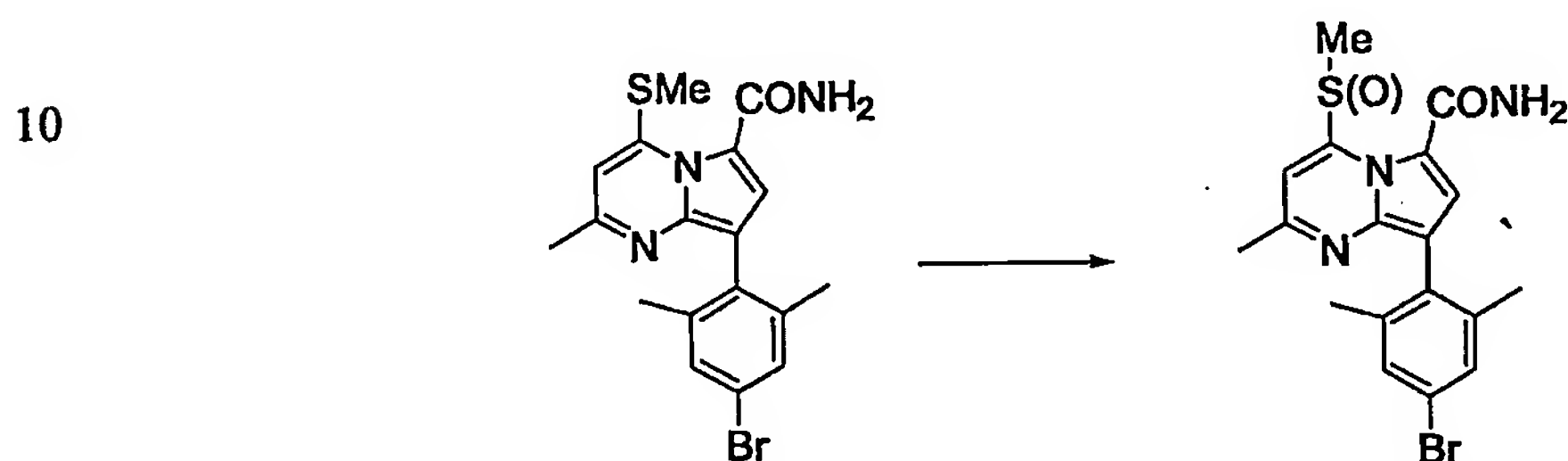
(2) A mixture of 8-(4-bromo-2,6-dimethyl-phenyl)-4-mercapto-2-methyl-pyrrolo[1,2-*a*]pyrimidine-6-carbonitrile (7.50 g), MeI (12.5 mL) in 1 M NaOH aqueous solution (100 mL) was stirred at room temperature for 1 h. The reaction mixture was extracted with ethyl acetate. The organic layer was washed with brine, dried over anhydrous sodium sulfate and filtered. The filtrate was concentrated under reduced pressure to give crude 8-(4-bromo-2,6-dimethyl-phenyl)-2-methyl-4-methylsulfanyl-pyrrolo[1,2-*a*]pyrimidine-6-carbonitrile (5.75 g). This product was used in the next step without further purification.

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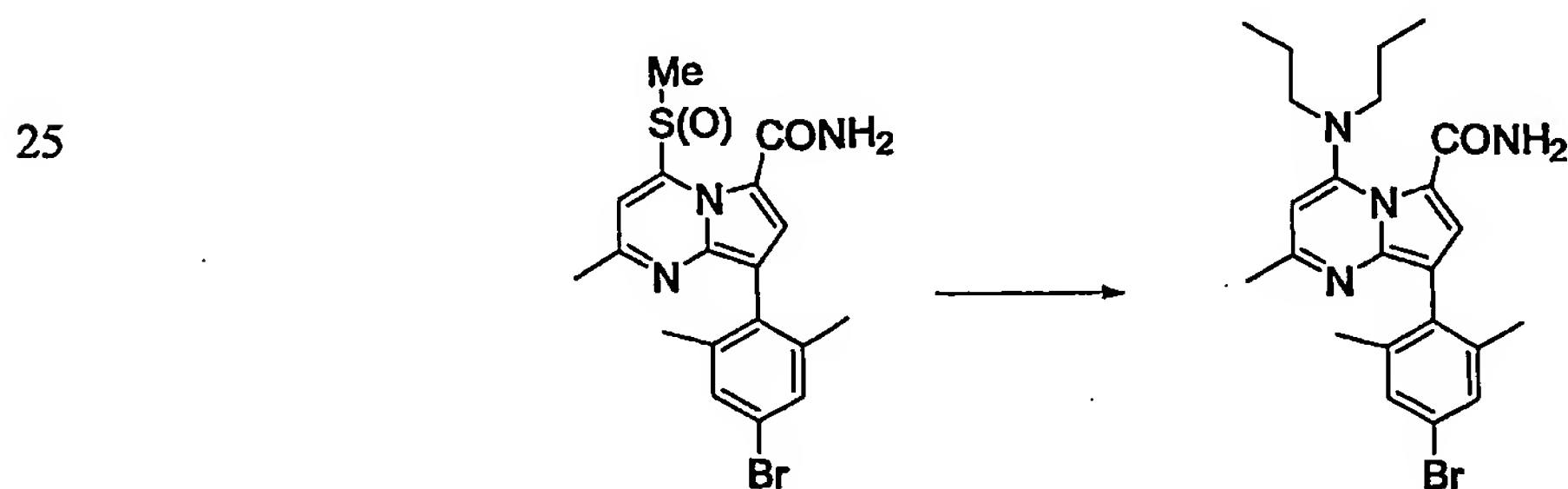


(3) 8-(4-bromo-2,6-dimethyl-phenyl)-2-methyl-4-methylsulfanyl-pyrrolo[1,2-

a]pyrimidine-6-carbonitrile (5.70 g) was added into conc. H₂SO₄ (100 mL) and heated for 60 °C for 5 hours. The reaction mixture was cooled to room temperature, poured into ice-water and then 10% aqueous NaOH solution was added to make the aqueous mixture alkaline (pH = 8) and extracted with ethyl acetate. The organic layer was washed with brine, dried over anhydrous sodium sulfate and filtered. The filtrate was concentrated under reduced pressure and the residue was purified by a silica gel column chromatography (silica gel: Wako Gel (C200), eluent: ethyl acetate) to give 8-(4-bromo-2,6-dimethyl-phenyl)-2-methyl-4-methylsulfanyl-pyrrolo[1,2-*a*]pyrimidine-6-carboxylic acid amide (3.12 g).



15 (4) To a solution of Oxone (9.12g) in water (50 mL) was added a solution of 8-(4-bromo-2,6-dimethyl-phenyl)-2-methyl-4-methylsulfanyl-pyrrolo[1,2-*a*]pyrimidine-6-carboxylic acid amide (3.00 g) in ethanol (50 mL) in an ice-cooling bath. The reaction mixture was stirred under ice-cooling for 30 minutes, poured into water and extracted with ethyl acetate. The organic layer was washed with a saturated aqueous sodium hydrogencarbonate and brine, dried over anhydrous sodium sulfate and filtered. The filtrate was concentrated under reduced pressure and the residue was purified by a silica gel column chromatography (silica gel: Wako Gel (C200), eluent: ethyl acetate) to give 8-(4-bromo-2,6-dimethyl-phenyl)-4-methanesulfinyl-2-methyl-pyrrolo[1,2-*a*]pyrimidine-6-carboxylic acid amide (1.68 g).

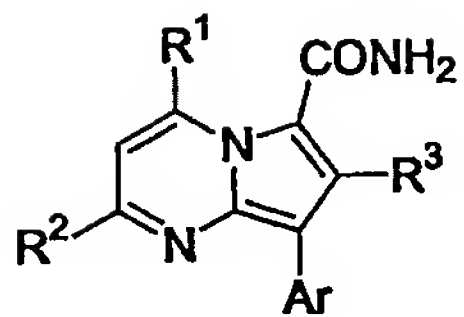


(5) A mixture of 8-(4-bromo-2,6-dimethyl-phenyl)-4-methanesulfinyl-2-methyl-pyrrolo[1,2-*a*]pyrimidine-6-carboxylic acid amide (100 mg), *N,N*-dipropylamine (48 mg) in ethanol (1 mL) was heated at reflux for 1 h. The reaction mixture was cooled to room temperature, poured into a saturated aqueous sodium hydrogencarbonate, and then extracted with ethyl acetate.

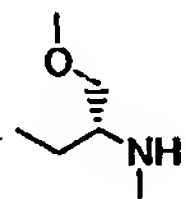
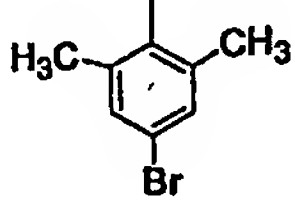
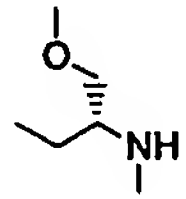
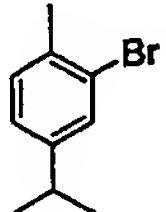
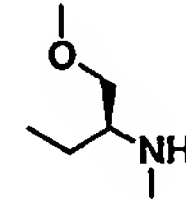
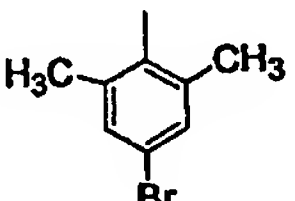
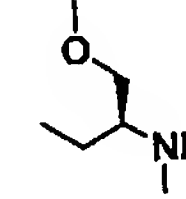
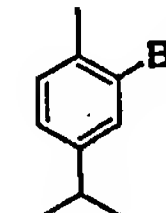
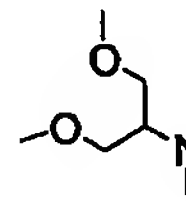
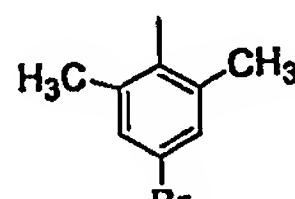
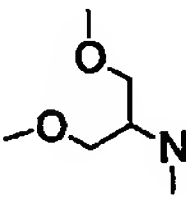
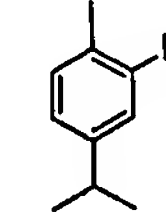
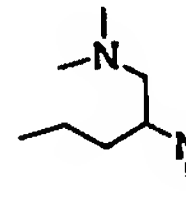
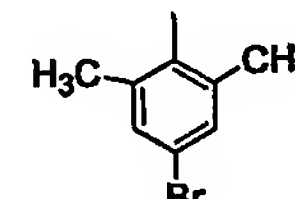
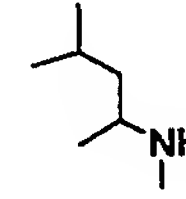
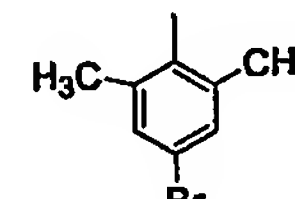
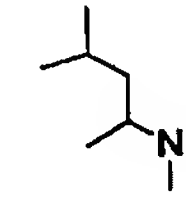
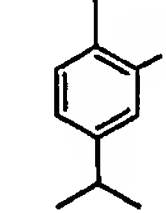
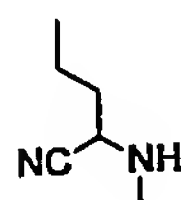
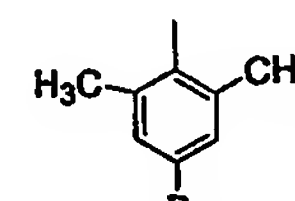
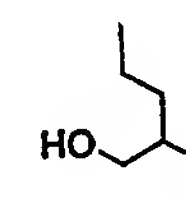
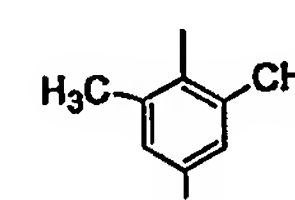
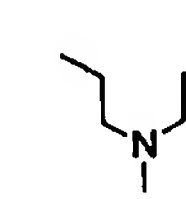
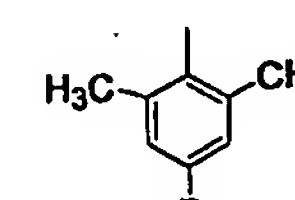
- 5 The organic layer was washed with brine, dried over anhydrous sodium sulfate and filtered. The filtrate was concentrated under reduced pressure and the residue was purified by a silica gel column chromatography (silica gel: Wako Gel (C200), eluent: hexane / ethyl acetate = 1 : 1) to give a solid. The solid was washed with a mixture of diisopropylether and ethyl acetate to give the title compound (50 mg).

10

Table 1 lists the compound obtained in Example 2 and compounds obtained by the similar procedure as in Example 2.

Table 1^{*1}

Com. No.	Ex. No.	R ¹	R ²	R ³	Ar	melting point (°C) (solvent for crystallization)
1-001	1		CH ₃	H		163-165 ^{*2} (EtOAc / EtOH)
1-002	1		CH ₃	H		195-197 ^{*2} (EtOAc / EtOH)
1-003	1		CH ₃	H		213-215 ^{*2} (EtOAc / EtOH)
1-004	1		CH ₃	H		204-206 ^{*2} (EtOAc / EtOH)
1-005	1		CH ₃	H		203-205 ^{*2} (EtOAc / EtOH)
1-006	1		CH ₃	H		180-182 ^{*2} (EtOAc / EtOH)
1-007	1		CH ₃	H		166-168 ^{*2} (EtOAc / EtOH)
1-008	1		CH ₃	H		175-177 ^{*2} (EtOAc / EtOH)

1-009	1		CH ₃	H		172-174 ^{*2} (EtOAc / EtOH)
1-010	1		CH ₃	H		160-162 ^{*2} (EtOAc / EtOH)
1-011	1		CH ₃	H		172-174 ^{*2} (EtOAc / EtOH)
1-012	1		CH ₃	H		166-168 ^{*2} (EtOAc / EtOH)
1-013	1		CH ₃	H		203-205 ^{*2} (EtOAc / EtOH)
1-014	1		CH ₃	H		188-190 ^{*2} (EtOAc / EtOH)
1-015	1		CH ₃	H		183-185 ^{*2} (EtOAc)
1-016	1		CH ₃	H		180-182 ^{*2*3}
1-017	1		CH ₃	H		163-165 ^{*2*3}
1-018	2		CH ₃	H		240-242 (EtOAc)
1-019	2		CH ₃	H		232-234 (decomp.) (EtOAc)
1-020	2		CH ₃	H		199-201 (EtOAc)

1-021	2		CH ₃	H		208-210 (EtOAc)
1-022	2		CH ₃	H		178-180 (EtOAc)
1-023	2		CH ₃	H		194-196 ^{*3}
1-024	2		CH ₃	H		amorphous
1-025	2		CH ₃	H		223-225 (EtOAc)
1-026	2		CH ₃	H		227-229 (EtOAc)
1-027	2		CH ₃	H		222-224 (EtOAc)
1-028	1		CH ₃	H		amorphous

*1: Com. No. = compound number, Ex. No. = example number, solvent for crystallization;

EtOAc = ethyl acetate, EtOH = ethanol

Analytical data of non-crystal compounds, diastereoisomers and optically active compounds are described below.

1-024:

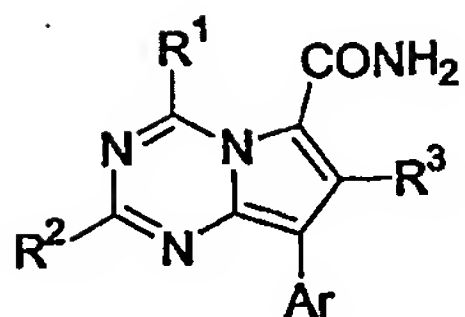
MS (Pos, ES): 442 (M + Na)⁺, 444 (M + Na + 2)⁺; NMR (300 MHz, CDCl₃) δ 2.04 (3 H, s), 2.09 (3 H, s), 2.58 (3 H, s), 3.17 (3 H, s), 5.54-5.66 (2 H, m), 7.26 (1 H, s), 7.31 (2 H, s), 7.59 (1 H, s)

1-028:

MS (Pos, ES): 486 ($M + 1$)⁺, 488 ($M + 3$)⁺, 508 ($M + Na$)⁺, 510 ($M + Na + 2$)⁺; NMR (300 MHz, CDCl₃) δ 0.98 (3 H, t, $J = 7.3$ Hz), 1.40-1.64 (2 H, m), 1.68-1.79 (2 H, m), 2.09 (3 H, s), 2.10 (3 H, s), 2.37 (3 H, s), 2.51 (1 H, dd, $J = 5.9, 14.4$ Hz), 2.65 (1 H, dd, $J = 7.4, 14.4$ Hz), 4.07-4.18 (1 H, m), 5.28-5.39 (1 H, br s), 5.48-5.58 (2 H, br s), 5.72-5.87 (1 H, br s), 5.89 (1 H, s), 7.22 (1 H, s), 7.26 (3 H, s), 10.75-10.92 (1 H, br s)

*2: HCl salt

*3: Crystallized on standing from the compound purified (silica gel column chromatography) and dried.

Table 2^{*1}

Com. No.	Ex. No.	R ¹	R ²	R ³	Ar	melting point (°C) (solvent for crystallization)
2-001	1		CH ₃	H		225-227 ^{*2} (EtOAc / IPE)
2-002	1		CH ₃	H		244-246 ^{*2} (EtOAc)
2-003	1		CH ₃	H		229-231 ^{*2} (EtOAc)
2-004	1		CH ₃	H		214-216 ^{*2} (EtOAc)
2-005	1		CH ₃	H		218-220 ^{*2} (EtOAc)
2-006	1		CH ₃	H		206-208 ^{*2} (decomp.) (EtOAc)

*1: Com. No. = compound number, Ex. No. = example number, solvent for crystallization;

EtOAc = ethyl acetate, IPE = diisopropylether

*2: HCl salt

Test Example [CRF receptor binding test]

Monkey amygdala membranes were used as a receptor preparation.

^{125}I -CRF was used as ^{125}I -labeled ligand.

Binding reaction using the ^{125}I -labeled ligand was carried out by the following method
5 described in The Journal of Neuroscience, 7, 88 (1987).

Preparation of receptor membranes:

Monkey amygdala was homogenized in 50 mM Tris-HCl buffer (pH 7.0) containing 10
mM MgCl_2 , 2 mM EDTA and centrifuged at 48,000 x g for 20 min, and the precipitate was
washed once with Tris-HCl buffer. The washed precipitate was suspended in 50 mM Tris-HCl
10 buffer (pH 7.0) containing 10 mM MgCl_2 , 2 mM EDTA, 0.1% bovine serum albumin and 100
kallikrein units/ml aprotinin, to obtain a membrane preparation.

CRF receptor binding test:

The membrane preparation (0.3 mg protein/ml), ^{125}I -CRF (0.2 nM) and a test drug were
reacted at 25°C for 2 hours. After completion of the reaction, the reaction mixture was filtered
15 by suction through a glass filter (GF/C) treated with 0.3% polyethylene imine, and the glass filter
was washed three times with phosphate-buffered saline containing 0.01% Triton X-100. After
the washing, the radioactivity of the filter paper was measured in a gamma counter.

The amount of ^{125}I -CRF bound when the reaction was carried out in the presence of 1
 μM CRF was taken as the degree of nonspecific binding of ^{125}I -CRF, and the difference between
20 the total degree of ^{125}I -CRF binding and the degree of nonspecific ^{125}I -CRF binding was taken as
the degree of specific ^{125}I -CRF binding. An inhibition curve was obtained by reacting a definite
concentration (0.2 nM) of ^{125}I -CRF with various concentrations of each test drug under the
conditions described above. A concentration of the test drug at which binding of ^{125}I -CRF is
inhibited by 50% (IC_{50}) was determined from the inhibition curve.

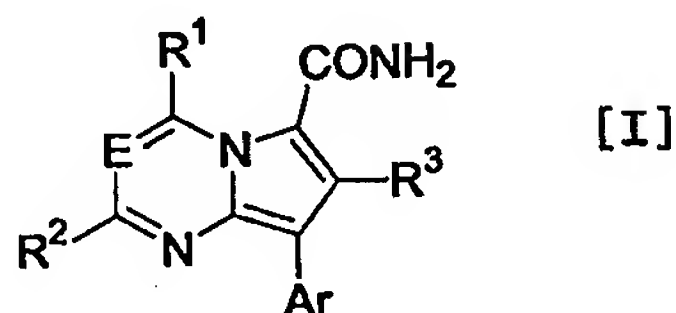
25 As a result, it was found that compounds 1-001, 1-002, 1-003, 1-004, 1-005, 1-006, 1-
007, 1-008, 1-009, 1-010, 1-011, 1-012, 1-013, 1-016, 1-017, 1-018, 1-019, 1-022, 1-027, 2-001,
2-002, 2-003, 2-004, 2-005 and 2-006 can be exemplified as typical compounds having an IC_{50}
value of 100 nM or less.

[EFFECT OF THE INVENTION]

According to the present invention, compounds having a high affinity for CRF receptors have been provided. These compounds are effective against diseases in which CRF is
5 considered to be involved, such as depression, anxiety, Alzheimer's disease, Parkinson's disease, Huntington's chorea, eating disorder, hypertension, gastral diseases, drug dependence, epilepsy, cerebral infarction, cerebral ischemia, cerebral edema, cephalic external wound, inflammation, immunity-related diseases, alpecia, irritable bowel syndrome, sleep disorders, epilepsy, dermatitides, schizophrenia, etc.

WHAT IS CLAIMED IS:

1. A pyrrolopyrimidine or pyrrolotriazine derivative substituted with a carbamoyl group represented by the following formula [I]:



(wherein E is N or CH;

R^1 is $-OR^4$, $-S(O)_lR^4$ or $-NR^4R^5$;

- 10 R^4 and R^5 are the same or different, and independently hydrogen, C_{1-9} alkyl, C_{3-7} cycloalkyl, C_{3-7} cycloalkyl- C_{1-6} alkyl, di(C_{3-7} cycloalkyl)- C_{1-6} alkyl, C_{1-6} alkoxy- C_{1-6} alkyl, di(C_{1-6} alkoxy)- C_{1-6} alkyl, hydroxy- C_{1-6} alkyl, cyano- C_{1-6} alkyl, carbamoyl- C_{1-6} alkyl or di(C_{1-6} alkyl)amino- C_{2-6} alkyl; or R^4 and R^5 are taken together to form $-(CH_2)_m-A-(CH_2)_n-$ wherein A is methylene, oxygen, sulfur, NR^6 or CHR^7 , m is an integer selected from 1, 2, 3 and 4, n is an
- 15 integer selected from 0, 1 and 2, wherein R^6 is hydrogen, C_{1-6} alkyl or C_{3-7} cycloalkyl, R^7 is hydroxy, hydroxy- C_{1-6} alkyl, cyano or cyano- C_{1-6} alkyl;

R^2 is hydrogen or C_{1-6} alkyl;

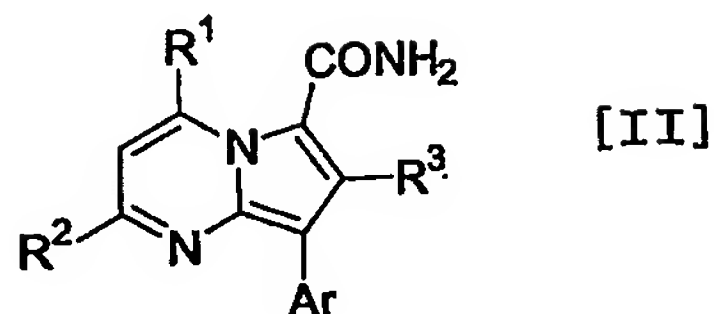
R^3 is hydrogen or C_{1-6} alkyl;

l is an interger selected from 0, 1 and 2;

- 20 Ar is aryl or heteroaryl which aryl or heteroaryl is unsubstituted or substituted with 1 or more substituents, which are the same or different, selected from the group consisting of halogen, C_{1-6} alkyl, C_{3-7} cycloalkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{1-6} alkoxy, C_{1-6} alkylthio, cyano, trifluoromethyl, trifluoromethoxy, difluoromethoxy, fluoromethoxy and $-N(R^8)R^9$, wherein R^8 and R^9 are the same or different, and independently are hydrogen or C_{1-6} alkyl)
- 25 , individual isomers thereof or racemic or non-racemic mixtures of isomers thereof, or pharmaceutically acceptable salts and hydrates thereof.

2. The pyrrolopyrimidine derivative substituted with a carbamoyl group according to claim

1 represented by the following formula [II]:



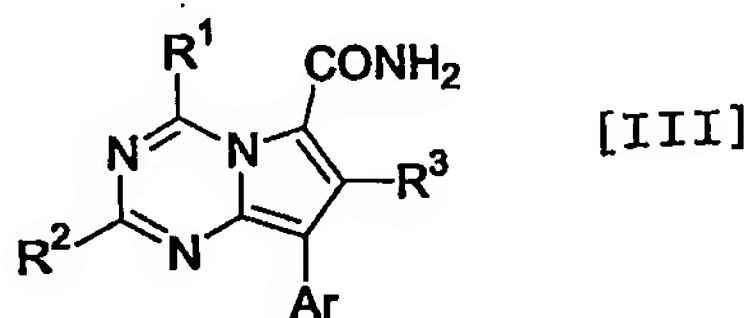
5

(wherein R^1 , R^2 , R^3 and Ar are as defined in claim 1), individual isomers thereof or racemic or non-racemic mixtures of isomers thereof, or pharmaceutically acceptable salts and hydrates thereof.

- 10 3. The pyrrolopyrimidine derivative substituted with a carbamoyl group according to claim 2 represented by the formula [II], wherein R^1 is $-NR^4R^5$; R^4 and R^5 are the same or different, and independently hydrogen, C_{1-9} alkyl, C_{3-7} cycloalkyl, C_{3-7} cycloalkyl- C_{1-6} alkyl, di(C_{3-7} cycloalkyl)- C_{1-6} alkyl, C_{1-6} alkoxy- C_{1-6} alkyl, di(C_{1-6} alkoxy)- C_{1-6} alkyl; R^2 is C_{1-6} alkyl; Ar is phenyl which phenyl is substituted with two or three substituents, which are the same or different, selected from
- 15 the group consisting of halogen, C_{1-3} alkyl, C_{1-3} alkoxy, C_{1-3} alkylthio, trifluoromethyl, trifluoromethoxy and $-N(R^8)R^9$ (wherein R^8 and R^9 are the same or different, and independently are hydrogen or C_{1-3} alkyl); R^3 is as defined in claim 1, individual isomers thereof or racemic or non-racemic mixtures of isomers thereof, or pharmaceutically acceptable salts and hydrates thereof.

20

4. The pyrrolotriazine derivative substituted with a carbamoyl group according to claim 1 represented by the following formula [III]:



25

(wherein R^1 , R^2 , R^3 and Ar are as defined in claim 1), individual isomers thereof or racemic or non-racemic mixtures of isomers thereof, or pharmaceutically acceptable salts and hydrates thereof.

thereof.

5. The pyrrolotriazine derivative substituted with a carbamoyl group according to claim 4 represented by the formula [III], wherein R^1 is $-NR^4R^5$; R^4 and R^5 are the same or different, and
 5 independently hydrogen, C_{1-9} alkyl, C_{3-7} cycloalkyl, C_{3-7} cycloalkyl- C_{1-6} alkyl, di(C_{3-7} cycloalkyl)- C_{1-6} alkyl, C_{1-6} alkoxy- C_{1-6} alkyl, di(C_{1-6} alkoxy)- C_{1-6} alkyl; R^2 is C_{1-6} alkyl; Ar is phenyl which phenyl is substituted with two or three substituents, which are the same or different, selected from the group consisting of halogen, C_{1-3} alkyl, C_{1-3} alkoxy, C_{1-3} alkylthio, trifluoromethyl, trifluoromethoxy and $-N(R^8)R^9$ (wherein R^8 and R^9 are the same or different, and independently
 10 are hydrogen or C_{1-3} alkyl); R^3 is as defined in claim 1, individual isomers thereof or racemic or non-racemic mixtures of isomers thereof, or pharmaceutically acceptable salts and hydrates thereof.

6. An antagonist for CRF receptors, comprising a pyrrolopyrimidine or pyrrolotriazine
 15 derivative substituted with a carbamoyl group, a pharmaceutically acceptable salt thereof or its hydrate according to any one of claims 1 to 5, as an active ingredient.

7. Use of a pyrrolopyrimidine or pyrrolotriazine derivative substituted with a carbamoyl group, a pharmaceutically acceptable salt thereof or its hydrate according to any one of claim 1 to
 20 5, for the manufacture of an antagonist for CRF receptors.

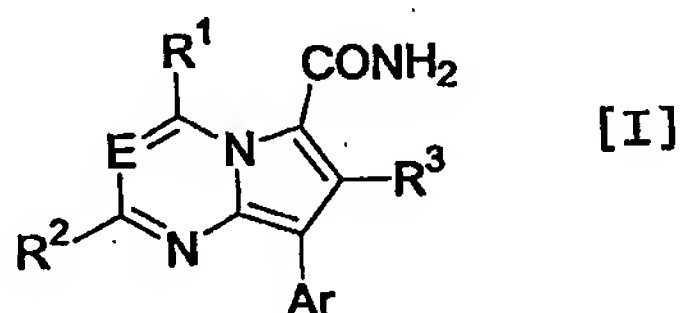
[ABSTRACT]

[PROBLEM TO BE SOLVED]

An object of the present invention is to provide an antagonist against CRF receptors which is effective as a therapeutic or prophylactic agent for diseases in which CRF is considered to be involved, such as depression, anxiety, Alzheimer's disease, Parkinson's disease, Huntington's chorea, eating disorder, hypertension, gastral diseases, drug dependence, epilepsy, cerebral infarction, cerebral ischemia, cerebral edema, cephalic external wound, inflammation, immunity-related diseases, alpecia, irritable bowel syndrome, sleep disorders, epilepsy, dermatitides, schizophrenia, etc.

[SOLUTION]

A pyrrolopyrimidine or pyrrolotriazine derivative substituted with a carbamoyl group represented by the following formula [I]:



has a high affinity for CRF receptors and is effective against diseases in which CRF is considered to be involved.

特願 2004-001311

出願人履歴情報

識別番号

[000002819]

1. 変更年月日

1990年 8月22日

[変更理由]

新規登録

住所

東京都豊島区高田3丁目24番1号

氏名

大正製薬株式会社